

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : C10L 1/18, C10M 145/08, 145/16, 145/12, 145/02		A1	(11) International Publication Number: WO 95/23200 (43) International Publication Date: 31 August 1995 (31.08.95)		
(21) International Application Number: PCT/EP95/00666		(81) Designated States: CA, CZ, FI, HU, JP, KR, NO, PL, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).			
(22) International Filing Date: 22 February 1995 (22.02.95)					
(30) Priority Data: 9403660.5 25 February 1994 (25.02.94) GB		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>			
(71) Applicant (for all designated States except US): EXXON CHEMICAL PATENTS INC. [US/US]; 1900 East Linden Avenue, Linden, NJ 07036 (US).					
(72) Inventors; and					
(75) Inventors/Applicants (for US only): BROWN, Gerald, Ivan [GB/GB]; 3 Orchard Gardens, West Challow, Wantage, Ox- fordshire OX12 9TL (GB). MORE, Iain [GB/GB]; 15 Lee Avenue, Abingdon, Oxfordshire OX14 3UT (GB). TACK, Robert, Dryden [GB/GB]; 42 Alexander Close, Abingdon, Oxfordshire OX14 1XB (GB). DAVIES, Brian, William [GB/GB]; Pound Furlong, London Road, Blewbury, Ox- fordshire OX11 9PD (GB). TOWE, Derek, Alan [GB/GB]; Croft Cottage, Ford Lane, East Hendred, Wantage, Oxford- shire O12 8JX (GB).					
(74) Agents: MANSELL, Keith, Rodney et al.; Exxon Chemical Limited, Exxon Chemical Technology Centre, P.O. Box 1, Abingdon, Oxfordshire OX13 6BB (GB).					
(54) Title: OIL COMPOSITIONS					
(57) Abstract					
Additives having certain hydrocarbyl groups improve the low temperature properties of hydrocarbon oils.					

BEST AVAILABLE COPY

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	GB	United Kingdom	MR	Mauritania
AU	Australia	GE	Georgia	MW	Malawi
BB	Barbados	GN	Guinea	NE	Niger
BE	Belgium	GR	Greece	NL	Netherlands
BF	Burkina Faso	HU	Hungary	NO	Norway
BG	Bulgaria	IE	Ireland	NZ	New Zealand
BJ	Benin	IT	Italy	PL	Poland
BR	Brazil	JP	Japan	PT	Portugal
BY	Belarus	KE	Kenya	RO	Romania
CA	Canada	KG	Kyrgyzstan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic of Korea	SD	Sudan
CG	Congo	KR	Republic of Korea	SE	Sweden
CH	Switzerland	KZ	Kazakhstan	SI	Slovenia
CI	Côte d'Ivoire	LJ	Liechtenstein	SK	Slovakia
CM	Cameroon	LK	Sri Lanka	SN	Senegal
CN	China	LU	Luxembourg	TD	Chad
CS	Czechoslovakia	LV	Latvia	TG	Togo
CZ	Czech Republic	MC	Monaco	TJ	Tajikistan
DE	Germany	MD	Republic of Moldova	TT	Trinidad and Tobago
DK	Denmark	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	US	United States of America
FI	Finland	MN	Mongolia	UZ	Uzbekistan
FR	France			VN	Viet Nam
GA	Gabon				

OIL COMPOSITIONS

This invention concerns oil compositions having improved low temperature properties, and additives imparting such properties to hydrocarbon oils.

5

The general problem of decreased hydrocarbon oil flowability at low temperatures is well recognised in the art. Hydrocarbon oils typically contain normal alkanes which precipitate from the bulk oil at and below oil cloud point temperature, forming wax crystals. These wax crystals modify the flow characteristics of the

10

hydrocarbon oil, eventually forming a spongy mass which entraps the bulk oil.

One well-recognised solution to this problem is the use of chemical additives to improve the flowability of hydrocarbon oil at temperatures below cloud point. This improvement may result from additive interaction with the forming wax crystals, for example by reducing crystal size, the smaller wax crystals appearing less likely to clog fine filters. Other additives inhibit wax crystallisation into platelets, instead causing the adoption of acicular crystal habits which pass through the filter pores more readily. Many such 'low temperature flow improving' additives have been described in the art.

20

However, such additives commonly exhibit the problem of having non-universal application across the range of hydrocarbon oil types. Typically, a particular additive will prove effective only in oils sharing certain physical characteristics, and will prove largely ineffective in other oils. It is a continual challenge to devise additives effective in a range of oils, and particularly in those oils hitherto regarded in the art as difficult to treat with conventional low-temperature flow improving additives.

30

Comb polymers generally have one or more long chain substituents pendant from a polymer backbone, said substituents being bonded either directly to the backbone or indirectly to the backbone via interposed atoms or groups. Comb polymers are discussed in "Comb-like Polymers, Structure and Properties" by N.A. Platé and V.P. Shibaev, published in J. Poly. Sci, Macromolecular Revs. 8, p. 117 to 253 (1974). A number of classes of comb polymer useful as low temperature flow improving additives have been described in the art.

UK Patent No. 1,469,016 describes comb polymers derived from C₆ to C₁₈ alkyl esters of unsaturated C₄ to C₈ dicarboxylic acids, with copolymers of di-n-alkyl

fumarates and vinyl acetate being preferred. Such comb polymers are shown to be effective as low temperature flow improvers only in fuel oils having high end points, i.e. final boiling points above 700°F (371°C).

5 European Patent Application No. 0,282,342 describes comb polymers derived from a C₂ to C₁₇ alpha-olefin or aromatic substituted olefin, and a mono- or di- C₈ to C₂₃ alkyl ester of certain unsaturated carboxylic acids. Such polymers are shown to be effective as low temperature flow improvers only in fuel oils having a relatively high final boiling point of above 360°C.

10 UK Patent No. 2,023,645 describes a three component additive combination for fuel oils, in which "Component B" is a comb polymer having hydrocarbyl substituents in the form of straight chain alkyl groups of 6 to 30 carbon atoms. Such additive combinations are shown to be effective as low temperature flow 15 improvers and as inhibitors of wax settling only in fuels having final boiling points of at least 361°C.

20 The known comb polymers hereinbefore described have not proved substantially advantageous in hydrocarbon oils lacking the physical characteristic of high final boiling point.

✓ WO 94/00386 discloses oil soluble ethylene polymers having, in addition to units derived from ethylene, units of the formulas

25 - CH₂ - CRR¹ - and - CH₂ - CRR² -,

30 wherein each R independently represents H or CH₃, and each R¹ and R² independently represents a group of the formula COOR³ or OOCR⁴, wherein R³ and R⁴ independently represent hydrocarbyl groups of, most preferably, at most 8 carbon atoms. Specifically-disclosed examples include ethylene vinyl n-octanoate (Example G) and ethylene vinyl n-heptanoate (Example F).

35 United States Patent No. 4,863,486 discloses a class of comb polymer surprisingly effective as a low temperature flow improver for distillate fuel oils which have a relatively narrow and/or low boiling range, and which are regarded as difficult to treat. Such fuel oils are described as having at least one of the following characteristics:

- (a) a final boiling point in the range of 340°C to 370°C
- (b) 20% and 90% distillation points differing by less than 100°C *(70-20%)*
- 5 (c) 90% distillation point and final boiling point differing by between 10°C and 25°C.

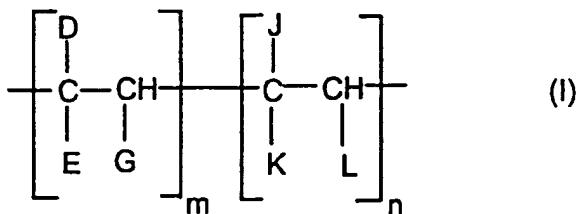
The comb polymers disclosed as effective in these fuels comprise at least 25 % wt of a monomer being the n-alkyl ester of a mono-ethylenically unsaturated C₃-C₈ 10 mono- or dicarboxylic acid, the average number of carbon atoms (hereinafter "average carbon number") in the n-alkyl groups being from 12 to 14 and the proportion of individual n-alkyl groups having more than 14, or less than 12, carbon atoms being strictly limited. Comb polymers having n-alkyl groups (i.e. hydrocarbyl substituents) not in accordance with this average carbon number 15 range are shown to be ineffective in such fuels.

Hydrocarbon oils - and particularly fuel oils - currently produced for winter use in many Scandinavian, North American and other cold regions typically have cloud 20 points of -10°C or below. Such oils often have a narrow and/or low boiling range, and also often have a low final boiling point. These oils are particularly difficult to treat with low-temperature flow improving additives. In particular, low cloud point oils appear, on cooling, to exhibit higher rates of wax crystallisation once the 25 cloud point is reached. This rapid deposition of solid wax appears to interfere with the action of conventional low temperature flow improvers.

25 In particular, we have found that the additives described in USP No. 4,863,486 as advantageous for distillate fuel oils having a relatively narrow and/or low boiling range are largely ineffective in oils having a cloud point no higher than -10°C, notwithstanding the distillation characteristics of these low cloud point oils. 30 Surprisingly, we have now found that the low temperature flowability of such low cloud point oils may be successfully improved through the use of comb polymers having hydrocarbyl substituents of average carbon number below 12. This improved oil flowability accordingly allows a mechanical system or device dependent, for normal operation, upon the flowability of these hydrocarbon oils to 35 remain operational at lower temperatures.

In a first aspect therefore, this invention provides an oil composition comprising a major proportion of hydrocarbon oil having a cloud point no higher than -10°C and

a minor proportion of an additive comprising a comb polymer containing units of the general formula (I)



5

wherein D represents COOR^{11} , OCOR^{11} or OR^{11} groups,

E represents H, CH_3 , D or R^{12} groups,

G represents H or D group,

J represents H, R^{12} or an aryl or heterocyclic group,

10 K represents H, COOR^{12} , OCOR^{12} , OR^{12} or COOH groups,

L represents H, R^{12} , COOR^{12} , OCOR^{12} or aryl groups, and

wherein R^{11} and R^{12} each represent hydrocarbyl substituents of average carbon number below 12 as measured over all units in the polymer, and
 15 wherein m and n represent mole ratios, their sum being 1 and m being finite and being up to and including 1 and n being from zero to less than 1, provided that E, G, J, K and L do not each represent H when D represents COOR^{11} or OCOR^{11} .

20 In a second aspect, this invention provides the use of an additive of the first aspect for improving the low temperature flow properties of a hydrocarbon oil having a cloud point no higher than -10°C .

25 In a third aspect, this invention provides the use of the oil composition of the first aspect in a mechanical system or device dependent, for normal operation, upon the flowability of hydrocarbon oil.

The invention is hereinafter described in more detail.

30 The Comb Polymer [of all aspects of the invention]

The hydrocarbyl substituents R^{11} and R^{12} are each of average carbon number below 12, preferably no higher than 11.75, more preferably no higher than 11, and

most preferably about 10. An average carbon number of 10 is especially advantageous.

5 In accordance with preferred embodiments of the invention, the hydrocarbyl substituents R¹¹ and R¹² each suitably have an average carbon number of above 8, and more suitably at least 9.

10 An individual unit (I) within the polymer may have hydrocarbyl substituents R¹¹ and R¹² which contain a different number of carbon atoms from those substituents in neighbouring units, provided that the relative proportions of different units within the polymer are such as to give the required average carbon number. Preferably, however, the individual units have hydrocarbyl substituents containing 15 approximately the same number of carbon atoms, such that the average carbon number for each of R¹¹ and R¹² is approximately equal to their carbon number in individual units. More preferably, the average carbon number for each of R¹¹ and R¹² is approximately equal to their carbon number in the predominant individual unit. Advantageously, all the individual units have substituents containing the same number of carbon atoms.

20 As used throughout this specification and not only in relation to the comb polymer, the term "hydrocarbyl" refers to a group being composed substantially or exclusively of carbon and hydrogen atoms such that the group is oleophilic in nature. Among these, there may be mentioned aliphatic, (e.g., alkyl or alkenyl), alicyclic (e.g., cycloalkyl or cycloalkenyl), aromatic, aliphatic- and alicyclic- 25 substituted aromatic, and aromatic-substituted aliphatic and alicyclic groups. Aliphatic groups are advantageously saturated.

30 Hydrocarbyl groups may contain substituents comprising hetero-atoms, provided they do not alter the oleophilic nature of the group. Examples include keto, halo, hydroxy, nitro, cyano, alkoxy, ester and acyl. If the hydrocarbyl group is substituted, a single (mono) substituent is preferred. Examples of substituted hydrocarbyl groups include 2-hydroxyethyl, 3-hydroxypropyl, 4-hydroxybutyl, 2-ketopropyl, ethoxyethyl, and propoxypropyl. The groups may also or alternatively contain hetero-atoms in a chain or ring otherwise composed of carbon atoms. 35 Suitable hetero-atoms include, for example, nitrogen, oxygen and sulfur.

The term "hydrocarbon" is used analogously throughout this specification.

In accordance with preferred embodiments of the invention, the individual hydrocarbyl substituents R¹¹ and R¹² are alkyl groups, and preferably n-alkyl groups such as n-octyl, n-nonyl, n-decyl, n-undecyl and n-dodecyl.

5 Preferably, D represents COOR¹¹ or OCOR¹¹ and E, G, J, K and L do not each represent H.

Particularly preferred embodiments are those in which R¹² represents a hydrocarbyl substituent containing from 1 to 6 carbon atoms and wherein R¹¹ 10 represents a hydrocarbyl substituent different from R¹² and of average carbon number below 12, and preferably above 8. In such embodiments, R¹² is advantageously subject to the preferences for average carbon number hereinbefore described. Thus, for example, the individual units of the polymer 15 preferably have R¹¹ substituents containing substantially the same number of carbon atoms.

The number average molecular weight of the polymers may, for example, be in the range of 1,000 to 120,000, preferably 1,000 to 50,000, more preferably 2,000 to 25,000, and most preferably 3,000 to 15,000 as measured by Vapour Phase 20 Osmometry (VPO).

Examples of particularly advantageous comb polymers include copolymers of one or more esters of an ethylenically unsaturated carboxylic acid such as maleic anhydride or fumaric acid and another ethylenically unsaturated monomer such as 25 an α -olefin or an unsaturated ester, for example vinyl acetate. It is preferred but not essential that equimolar amounts of the comonomers be used although molar proportions in the range of 2 to 1 and 1 to 2 are suitable. Examples of olefins that may be copolymerized with e.g. maleic anhydride include 1-octene, 1-decene, 1-dodecene and 1-tetradecene.

30 The copolymer may be esterified by any suitable technique and although preferred it is not essential that the maleic anhydride or fumaric acid be at least 50% esterified. Examples of alcohols which may be used include octan-1-ol, nonan-1-ol, decan-1-ol, undecan-1-ol, dodecan-1-ol and tetradecan-1-ol. The 35 alcohols may also include up to one methyl branch per chain, for example, 1-methyldecan-1-ol, 2-methyldecan-1-ol. The alcohol may be a mixture of normal and single methyl branched alcohols. Preferably, the alcohol contains only

normal alcohols. It is preferred to use pure alcohols rather than the commercially available alcohol mixtures.

Particularly preferred comb polymers are copolymers of alkyl fumarates and vinyl acetate made, for example, by solution copolymerizing an equimolar mixture of fumaric acid and vinyl acetate and reacting the resulting copolymer with the alcohol or mixture of alcohols, which are preferably straight chain alcohols. The particularly preferred fumarate comb polymers may, for example, have a number average molecular weight in the range of 1,000 to 100,000, preferably 1,000 to 10,000, more preferably 2,000 to 20,000, as measured by Vapour Phase Osmometry (VPO).

Other examples of particularly advantageous comb polymers are the copolymers of α -olefins, esterified copolymers of styrene and maleic anhydride, and esterified copolymers of styrene and fumaric acid. Further examples include those copolymers of esters of other ethylenically unsaturated carboxylic acids such as acrylic acid, methacrylic acid, itaconic acid and crotonic acid. Copolymers of these esters with vinyl esters of saturated carboxylic acids, in particular vinyl acetate and vinyl propionate, are especially suitable.

The comb polymers are generally prepared by polymerising the monomers in a solution of a hydrocarbon solvent such as heptane, benzene, cyclohexane, or white oil, at a temperature generally in the range of from 20°C to 150°C, and usually promoted with a peroxide or azo type catalyst such as benzoyl peroxide or azodiisobutyronitrile, under a blanket of an inert gas such as nitrogen or carbon dioxide in order to exclude oxygen. The polymer may be prepared under pressure in an autoclave, or by refluxing or other polymerisation methods known to the man skilled in the art.

Two or more comb polymers in accordance with this invention may be used, in combination, to advantageous effect.

The Additive [of all aspects of this invention]

The additive comprises the above-described comb polymer, or a mixture of such comb polymers, optionally in the form of a concentrate. In such a concentrate, the copolymer(s) may be dissolved in a solvent at a concentration within wide limits according to the needs and restrictions of the specific application, for example

from 1:90, such as 10:80, per cent (weight:weight). Examples of suitable solvents are hydrocarbons or oxygen-containing hydrocarbons such as kerosene, aromatic naphthas, and mineral lubricating oils.

- 5 The concentration of the additive in the oil may for example be in the range of 1 to 5,000 ppm of additive (active ingredient) by weight per weight of oil, for example 10 to 5,000 ppm such as 10 to 2000 ppm (active ingredient) by weight per weight of oil, preferably 25 to 500 ppm, more preferably 100 to 200 ppm.
- 10 The additive or additives should be soluble in the oil to the extent of at least 1000 ppm by weight of oil at ambient temperature. However, at least some of the additive may come out of solution near the cloud point of the oil in order to modify the wax crystals that form.
- 15 The one or more comb polymers in accordance with this invention may also be used in combination with co-additives, in particular conventional low temperature flow improvers, to advantageous effect. Preferably therefore, the additive of all aspects of the invention comprises one or more of the additional low temperature flow improvers hereinafter described.

20

Additional low temperature flow improvers:

Such co-additives may be selected from the following:

- 25 (i) a linear compound,
- ✓ (ii) an ethylene/unsaturated ester copolymer,
- 30 (iii) a polar, organic, nitrogen-containing wax crystal growth inhibitor
- (iv) a hydrocarbon polymer
- (v) a sulphur carboxy compound, and
- 35 (vi) a hydrocarbylated aromatic.

These additional low temperature flow improvers will now be discussed in more detail.

(i) **Linear Compounds**

Such compounds comprise a compound in which at least one substantially linear alkyl group having 10 to 30 carbon atoms is connected to a non-polymeric organic residue to provide at least one linear chain of atoms that includes the carbon atoms of said alkyl groups and one or more non-terminal oxygen atoms.

By "substantially linear" is meant that the alkyl group is preferably straight chain, but that essentially straight chain alkyl groups having a small degree of branching such as in the form of a single methyl group may be used.

Preferably, the compound has at least two of said alkyl groups when the linear chain may include the carbon atoms of more than one of said alkyl groups. When the compound has at least three of said alkyl groups, there may be more than one of such linear chains, which chains may overlap. The linear chain or chains may provide part of a linking group between any two such alkyl groups in the compound.

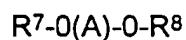
The oxygen atom or atoms are preferably directly interposed between carbon atoms in the chain and may, for example, be provided in the form of a mono- or poly-oxyalkylene group, said oxyalkylene group preferably having 2 to 4 carbon atoms, examples being oxyethylene and oxypropylene.

As indicated the chain or chains include carbon and oxygen atoms. They may also include other hetero-atoms such as nitrogen atoms.

The compound may be an ester where the alkyl groups are connected to the remainder of the compound as -O-CO n alkyl, or -CO-O n alkyl groups, in the former the alkyl groups being derived from an acid and the remainder of the compound being derived from a polyhydric alcohol and in the latter the alkyl groups being derived from an alcohol and the remainder of the compound being derived from a polycarboxylic acid. Also, the compound may be an ester where the alkyl groups are connected to the remainder of the compound as —O—n—alkyl groups. The compound may be both an ester and an ether or it may contain different ester groups.

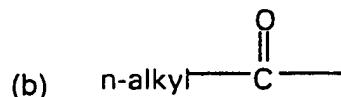
Examples are polyoxyalkylene esters, ethers, ester/ethers and mixtures thereof, particularly those containing at least one, preferably at least two C₁₀ to C₃₀ linear saturated alkyl groups and a polyoxyalkylene glycol group of molecular weight up to 5,000 preferably 200 to 5,000, the alkyl group in said polyoxyalkylene glycol containing from 1 to 4 carbon atoms. These materials form the subject of European Patent Publication 0 061 895 A2. Other such additives are described in United States Patent 4 491 455.

10 The preferred esters, ethers or ester/ethers which may be used may be structurally depicted by the formula

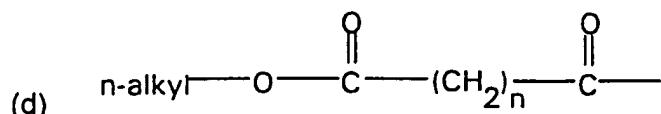
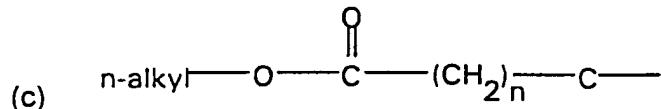


15 where R⁷ and R⁸ are the same or different and may be

(a) n-alkyl



20



25 n being, for example, 1 to 30, the alkyl group being linear and saturated and containing 10 to 30 carbon atoms, and A representing the polyalkylene segment of the glycol in which the alkylene group has 1 to 4 carbon atoms, such as a polyoxymethylene, polyoxyethylene or polyoxytrimethylene moiety which is substantially linear; some degree of branching with lower alkyl side chains (such as in polyoxypropylene glycol) may be present but it is preferred that the glycol is substantially linear. A may also contain nitrogen.

30

Examples of suitable glycols are substantially linear polyethylene glycols (PEG) and polypropylene glycols (PPG) having a molecular weight of about 100 to 5,000, preferably about 200 to 2,000. Esters are preferred and fatty acids containing from 10 to 30 carbon atoms are useful for reacting with the glycols to form the ester additives, it being preferred to use a C₁₈ to C₂₄ fatty acid, especially behenic acid. The esters may also be prepared by esterifying polyethoxylated fatty acids or polyethoxylated alcohols.

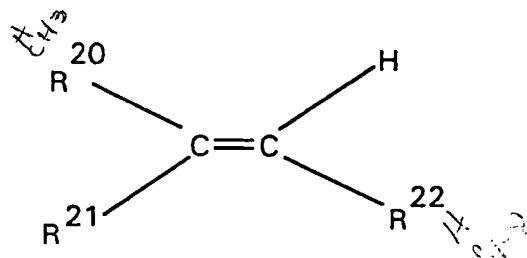
5 Polyoxyalkylene diesters, diethers, ether/esters and mixtures thereof are suitable
10 as additives, diesters being preferred for use in narrow boiling distillates when minor amounts of monoethers and monoesters (which are often formed in the manufacturing process) may also be present. It is important for additive performance that a major amount of the dialkyl compound is present. In particular, stearic or behenic diesters of polyethylene glycol, polypropylene glycol or polyethylene/polypropylene glycol mixtures are preferred.

15 Other examples of polyoxyalkylene compounds are those described in Japanese Patent Publication Nos. 2-51477 and 3-34790 (both Sanyo), and the esterified alkoxylated amines described in EP-A-117,108 and EP-A-326,356 (both Nippon Oil and Fats).

(ii) Ethylene/Unsaturated Ester Copolymers

20 Ethylene copolymer flow improvers have a polymethylene backbone divided into segments by oxyhydrocarbon side chains, i.e. ethylene unsaturated ester copolymer flow improvers. The unsaturated monomers copolymerisable with ethylene to form the copolymers include unsaturated mono and diesters of the general formula:

30



wherein R²⁰ represents hydrogen or a methyl group; R²¹ represents a -OOCR²³ or -COOR²³ group wherein R²³ represents hydrogen or a C₁ to C₈, straight or branched chain alkyl group, provided that R²³ does not

represent hydrogen when R²¹ represents -COOR²³; and R²² is hydrogen or -COOR²³.

The monomer, when R²⁰ and R²² are hydrogen and R²¹ is -OOCR²³, includes 5 vinyl alcohol esters of C₁ to C₈, preferably C₁ to C₅, monocarboxylic acids, and most preferably C₂ to C₅ monocarboxylic acids. Examples of vinyl esters which may be copolymerised with ethylene include vinyl acetate, vinyl propionate and vinyl butyrate or isobutyrate, vinyl acetate and vinyl propionate being preferred. 10 Preferably, the copolymers contain from 5 to 40 wt% of the vinyl ester, more preferably from 10 to 35 wt% vinyl ester. They may also be in the form of mixtures of two copolymers such as those described in US Patent 3,961,916. Preferably, number average molecular weight of the copolymer as measured by vapour phase osmometry is 1,000 to 10,000, more preferably 1,000 to 5,000. If desired, the 15 copolymers may be derived from additional comonomers, e.g. they may be terpolymers or tetrapolymers or higher polymers, for example where the additional comonomer is isobutylene or diisobutylene.

Such copolymers may also be made by transesterification, or by hydrolysis and 20 re-esterification, of an ethylene unsaturated ester copolymer to give a different ethylene unsaturated ester copolymer. For example, ethylene vinyl hexanoate and ethylene vinyl octanoate copolymers may be made in this way, e.g. from an ethylene vinyl acetate copolymer.

(iii) Polar Organic, Nitrogen-containing Compounds

25 The oil-soluble polar nitrogen compound is either ionic or non-ionic and is capable of acting as a wax crystal growth inhibitor in fuels. It comprises for example one or more of the compounds (a) to (c) as follows:

30 (a) An amine salt and/or amide formed by reacting at least one molar proportion of a hydrocarbyl substituted amine with a molar proportion of a hydrocarbyl acid having 1 to 4 carboxylic acid groups or its anhydride.

35 Ester/amides may be used containing 30 to 300, preferably 50 to 150 total carbon atoms. These nitrogen compounds are described in US Patent 4 211 534. Suitable amines are usually long chain C₁₂ to C₄₀ primary, secondary, tertiary or quaternary amines or mixtures thereof but shorter chain amines may be used provided the resulting nitrogen compound is oil

soluble and therefore normally contains about 30 to 300 total carbon atoms. The nitrogen compound preferably contains at least one straight chain C₈ to C₄₀, preferably C₁₄ to C₂₄, alkyl segment.

5 Suitable amines include primary, secondary, tertiary or quaternary, but preferably are secondary. Tertiary and quaternary amines can only form amine salts. Examples of amines include tetradecyl amine, cocoamine, and hydrogenated tallow amine. Examples of secondary amines include dioctacedyl amine and methyl-behenyl amine. Amine mixtures are also
10 suitable such as those derived from natural materials. A preferred amine is a secondary hydrogenated tallow amine of the formula HNR⁹R¹⁰ wherein R⁹ and R¹⁰ are alkyl groups derived from hydrogenated tallow fat composed of approximately 4% C₁₄, 31% C₁₆, 59% C₁₈.

15 Examples of suitable carboxylic acids and their anhydrides for preparing the nitrogen compounds include cyclohexane 1,2 dicarboxylic acid, cyclohexene 1,2 dicarboxylic acid, cyclopentane 1,2 dicarboxylic acid and naphthalene dicarboxylic acid, and 1,4-dicarboxylic acids including dialkyl
20 spirobislactone. Generally, these acids have about 5 to 13 carbon atoms in the cyclic moiety. Preferred acids useful in the present invention are benzene dicarboxylic acids such as phthalic acid, isophthalic acid, and terephthalic acid. Phthalic acid or its anhydride is particularly preferred. The particularly preferred compound is the amide-amine salt formed by reacting 1 molar portion of phthalic anhydride with 2 molar portions of
25 dihydrogenated tallow amine. Another preferred compound is the diamide formed by dehydrating this amide-amine salt.

30 Other examples are long chain alkyl or alkylene substituted dicarboxylic acid derivatives such as amine salts of monoamides of substituted succinic acids, examples of which are known in the art and described in US-A-4 147 520, for example. Suitable amines may be those described above.

35 Other examples are condensates such as described in EP-A-327,423.

35 (b) A chemical compound comprising or including a cyclic ring system, the compound carrying at least two substituents of the general formula (I) below on the ring system

$-A-NR^{13}R^{14}$

(I)

5 where A is an aliphatic hydrocarbyl group that is optionally interrupted by one or more hetero atoms and that is straight chain or branched, and R¹³ and R¹⁴ are the same or different and each is independently a hydrocarbyl group containing 9 to 40 carbon atoms optionally interrupted by one or more hetero atoms, the substituents being the same or different and the compound optionally being in the form of a salt thereof.

10 Preferably, A has from 1 to 20 carbon atoms and is preferably a methylene or polymethylene group.

15 Examples of substituted hydrocarbyl groups include 2-hydroxyethyl, 3-hydroxypropyl, 4-hydroxybutyl, 2-ketopropyl, ethoxyethyl, and propoxypropyl. The groups may also or alternatively contain atoms other than carbon in a chain or ring otherwise composed of carbon atoms. Suitable hetero atoms include, for example, nitrogen, sulphur, and, preferably, oxygen.

20 The cyclic ring system may include homocyclic, heterocyclic, or fused polycyclic assemblies, or a system where two or more such cyclic assemblies are joined to one another and in which the cyclic assemblies may be the same or different. Where there are two or more such cyclic assemblies, the substituents of the general formula (I) may be on the same or different assemblies, preferably on the same assembly. Preferably, the or each cyclic assembly is aromatic, more preferably a benzene ring. Most preferably, the cyclic ring system is a single benzene ring when it is preferred that the substituents are in the ortho or meta positions, which benzene ring may be optionally further substituted.

25 The ring atoms in the cyclic assembly or assemblies are preferably carbon atoms but may for example include one or more ring N, S or O atom, in which case or cases the compound is a heterocyclic compound.

Examples of such polycyclic assemblies include:

- (i) condensed benzene structures such as naphthalene, anthracene, phenanthrene, and pyrene;
- 5 (ii) condensed ring structures where none of or not all of the rings are benzene such as azulene, indene, hydroindene, fluorene, and diphenylene oxide;
- 10 (iii) rings joined "end-on" such as diphenyl;
- (iv) heterocyclic compounds such as quinoline, indole, 2:3 dihydroindole, benzofuran, coumarin, isocoumarin, benzothiophen, carbazole and thiadiphenylamine;
- 15 (v) non-aromatic or partially saturated ring systems such as decalin (i.e. decahydronaphthalene), α -pinene, cardinene, and bornylene; and
- 20 (vi) three-dimensional structures such as norbornene, bicycloheptane (i.e. norbornane), bicyclooctane, and bicyclooctene.

Each hydrocarbyl group constituting R^{13} and R^{14} in the invention (Formula I) may for example be an alkyl or alkylene group or a mono- or poly-alkoxyalkyl group. Preferably, each hydrocarbyl group is a straight chain alkyl group. The number of carbon atoms in each hydrocarbyl group is preferably 16 to 40, more preferably 16 to 24.

Also, it is preferred that the cyclic system is substituted with only two substituents of the general formula (I) and that A is a methylene group.

30 Examples of salts of the chemical compounds are the acetate and the hydrochloride.

35 The compounds may conveniently be made by reducing the corresponding amide which may be made by reacting a secondary amine with the appropriate acid chloride; and

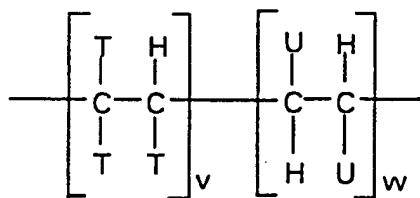
(c) A condensate of long chain primary or secondary amine with a carboxylic acid-containing polymer.

5 Specific examples include polymers such as described in GB-A-2,121,807,
 10 FR-A-2,592,387 and DE-A-3,941,561; and also esters of telemer acid and
 15 alkanoloamines such as described in US-A-4,639,256; a long chain
 epoxide/amine reaction product which may optionally be further reacted
 with a polycarboxylic acid; and the reaction product of an amine containing
 a branched carboxylic acid ester, an epoxide and a mono-carboxylic acid
 20 polyester such as described in US-A-4,631,071.

(iv) **Hydrocarbon Polymers**

Examples are those represented by the following general formula

15



where T = H or R¹⁵

U = H, T or aryl

20 R¹⁵ = C₁ to C₃₀ hydrocarbyl

and v and w represent mole ratios, v being within the range 1.0 to 0.0, w being within the range 0.0 to 1.0.

25 These polymers may be made directly from ethylenically unsaturated monomers or indirectly by hydrogenating the polymer made from monomers such as isoprene and butadiene.

30 Preferred hydrocarbon polymers are copolymers of ethylene and at least one α -olefin, having a number average molecular weight of at least 30,000. Preferably the α -olefin has at most 20 carbon atoms. Examples of such olefins are propylene, 1-butene, isobutene, n-octene-1, iso-octene-1, n-decene-1, and n-dodecene-1. The copolymer may also comprise small amounts, e.g. up to 10% by weight of other copolymerizable monomers, for example olefins other than α -

olefins, and non-conjugated dienes. The preferred copolymer is an ethylene-propylene copolymer. It is within the scope of the invention to include two or more different ethylene- α -olefin copolymers of this type.

5 The number average molecular weight of the ethylene- α -olefin copolymer is, as indicated above, at least 30,000, as measured by gel permeation chromatography (GPC) relative to polystyrene standards, advantageously at least 60,000 and preferably at least 80,000. Functionally no upper limit arises but difficulties of mixing result from increased viscosity at molecular weights above about 150,000, 10 and preferred molecular weight ranges are from 60,000 and 80,000 to 120,000.

15 Advantageously, the copolymer has a molar ethylene content between 50 and 85 per cent. More advantageously, the ethylene content is within the range of from 57 to 80%, and preferably it is in the range from 58 to 73%, more preferably from 62 to 71%, and most preferably 65 to 70%.

20 Preferred ethylene- α -olefin copolymers are ethylene-propylene copolymers with a molar ethylene content of from 62 to 71% and a number average molecular weight in the range 60,000 to 120,000, especially preferred copolymers are ethylene-propylene copolymers with an ethylene content of from 62 to 71% and a molecular weight from 80,000 to 100,000.

25 The copolymers may be prepared by any of the methods known in the art, for example using a Ziegler type catalyst. Advantageously, the polymers are substantially amorphous, since highly crystalline polymers are relatively insoluble in fuel oil at low temperatures.

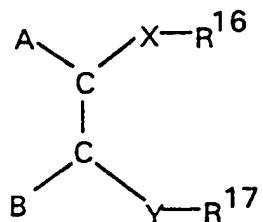
30 The additive composition may also comprise a further ethylene- α -olefin copolymer, advantageously with a number average molecular weight of at most 7500, advantageously from 1,000 to 6,000, and preferably from 2,000 to 5,000, as measured by vapour phase osmometry. Appropriate α -olefins are as given above, or styrene, with propylene again being preferred. Advantageously the ethylene content is from 60 to 77 molar per cent although for ethylene-propylene copolymers up to 86 molar per cent by weight ethylene may be employed with 35 advantage.

Examples of hydrocarbon polymers are described in WO-A-9 111 488.

(v) Sulphur Carboxy Compounds

Examples are those described in EP-A-0,261,957 which describes the use of compounds of the general formula

5



in which $-Y-R^{17}$ is $SO_3(-)(+)-NR\frac{18}{3}R^{17}$, $-SO_3(-)(+)-HNR\frac{18}{2}R^{17}$,

10 $-SO_3(-)(+)-H_2NR^{18}R^{17}$, $-SO_3(-)(+)-H_3NR^{17}$,
 $-SO_2NR^{18}R^{17}$, $-SO_3(-)(+)-H_3NR^{17}$;

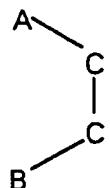
$-X-R^{16}$ is $-Y-R^{17}$ or $-CONR^{18}R^{16}$,
 $-CO_2(-)(+)-NR\frac{18}{3}R^{16}$, $-CO_2(-)(+)-HNR\frac{18}{2}R^{16}$,

15 $-R^{19}-COOR^{16}$, $-NR^{18}COR^{16}$,
 $-R^{19}OR^{16}$, $-R^{19}OCOR^{16}$, $-R^{19}, R^{16}$,
 $-N(COR^{18})R^{16}$ or $Z(-)(+)-NR\frac{18}{3}R^{16}$;
 $-Z(-)$ is $SO_3(-)$ or $-CO_2(-)$;

20 R^{16} and R^{17} are alkyl, alkoxyalkyl or polyalkoxyalkyl containing at least 10 carbon atoms in the main chain;

R^{18} is hydrocarbyl and each R^{18} may be the same or different and R^{19} is absent or is C_1 to C_5 alkylene and in

25



the carbon-carbon (C-C) bond is either a) ethylenically unsaturated when A and B may be alkyl, alkenyl or substituted hydrocarbyl groups or b) part of a cyclic

structure which may be aromatic, polynuclear aromatic or cyclo-aliphatic, it is preferred that X-R₁₆ and Y-R₁₇ between them contain at least three alkyl, alkoxyalkyl or polyalkoxyalkyl groups.

5 (vi) **Hydrocarbylated-Aromatics**

These materials are condensates comprising aromatic and hydrocarbyl parts. The aromatic part is conveniently an aromatic hydrocarbon which may be unsubstituted or substituted with, for example, non-hydrocarbon substituents. 10 Such an aromatic hydrocarbon preferably contains a maximum of these substituent groups and/or three condensed rings, and is preferably naphthalene. The hydrocarbyl part is a hydrogen and carbon containing part connected to the rest of the molecule by a carbon atom. It may be saturated or unsaturated, and straight or branched, and may contain one or more hetero-atoms provided they do 15 not substantially affect the hydrocarbyl nature of the part. Preferably the hydrocarbyl part is an alkyl part, conveniently having more than 8 carbon atoms. The molecular weight of such condensates may, for example, be in the range of 2,000 to 200,000 such as 2,000 to 20,000, preferably 2,000 to 8,000. Examples 20 are known in the art, primarily as lube oil pour depressants and dewaxing aids and they may, for example, be made by condensing a halogenated wax with an aromatic hydrocarbon. More specifically, the condensation may be a Friedel- Crafts condensation where the halogenated wax contains 15 to 60, e.g. 16 to 50, carbon atoms, has a melting point of about 200 to 400°C and has been 25 chlorinated to 5 to 25 wt% chlorine, e.g. 10 to 18 wt%. Another way of making similar condensates may be from olefins and the aromatic hydrocarbons.

Multicomponent additive systems may be used and the ratios of additives to be used will depend on the fuel to be treated.

30 In general, the additive of this invention is also suitable for use in hydrocarbon oils comprising other co-additives known in the art to impart beneficial properties to such oils. Amongst such other co-additives are the ashless dispersants described in numerous patent specifications, such as EP-A-0 482 253. Further examples include macrocyclic ashless dispersants, cetane improvers, polymers of 35 monoolefins, metallic-based combustion improvers such as ferrocene, corrosion inhibitors, anti-oxidants, reoadorants, antiwear additives, various emissions-reducing agents, and those hereinafter described in relation to the hydrocarbon oil.

The addition of these other co-additives may be simultaneous with that of the additives hereinbefore described; for example, the additive of the invention may additionally comprise one or more of the desired other co-additives. Alternatively, 5 the other co-additives may be added independent of the additive of the invention.

The Hydrocarbon Oil [of all aspects of this invention]

10 The hydrocarbon oil has a cloud point no higher than -10°C. In a preferred embodiment of this invention, the oil has a cloud point no higher than -12°C and in a more preferred embodiment, no higher than -14°C. Hydrocarbon oils having cloud points no higher than -20°C have proved particularly advantageous.

15 In this specification 'cloud point' refers to the physical characteristic determined in accordance with I.S.O. 3015 standard test procedure.

20 In general, the hydrocarbon oils useful in this invention may possess any distillation characteristics. However, in practice oils having the requisite low cloud point typically have relatively low final boiling points. Oils particularly suitable for this invention are therefore those having a final boiling point no higher than 370°, preferably no higher than 360°C, as measured by ASTM D-86.

25 Similarly, oils having the pre-requisite low cloud points typically exhibit relatively narrow boiling range. Such oils are particularly suitable for the invention and have 20% and 90% distillation points differing by less than 100°C, as measured by ASTM D-86.

30 Hydrocarbon oils having both relatively low final boiling points and relatively narrow boiling ranges, in addition to the pre-requisite low cloud points are especially suitable for this invention.

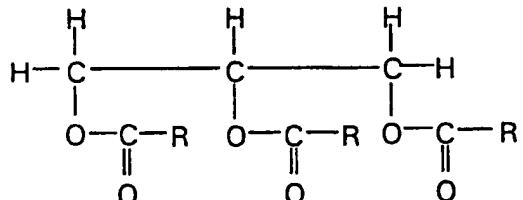
The hydrocarbon oil may be a crude oil, ie. an oil obtained directly from drilling and before refining.

35 The hydrocarbon oil may be a lubricating oil which may be an animal, vegetable or mineral oil, such as petroleum oil fractions ranging from naphthas or spindle oil to lubricating oil grades, castor oil, fish oils or oxidised mineral oil.

The hydrocarbon oil may preferably be a petroleum-based fuel oil, suitably a middle distillate fuel oil, ie, a fuel oil obtained in refining crude oil as the fraction between the lighter kerosene and jet fuels fraction and the heavier fuel oil fraction. The petroleum-based fuel oil can comprise atmospheric distillate or vacuum distillate, or cracked gas oil or a blend in any proportion of straight run and thermally and/or catalytically cracked distillates. The most common petroleum-based fuel oils are kerosene, jet fuels, diesel fuel, heating oils and heavy fuel oils.

These fuel oils may have a sulphur concentration of 0.2% by weight or less based on the weight of the fuel oil. Preferably, the sulphur concentration is 0.05% by weight or less, more preferably 0.01% by weight or less. The art describes methods for reducing the sulphur concentration of middle distillate fuel oils, such methods including solvent extraction, sulphuric acid treatment, and hydrodesulphurisation.

The hydrocarbon oil may be an oil derived from animal or vegetable material. Generally, such oils contain glycerides of a number of acids, the number and kind varying with the source of the oil. Vegetable oils are mainly tricyclerides of monocarboxylic acids, eg. acids containing 10-25 carbon atoms and of the formula



where R is an aliphatic radical of 10 to 25 carbon atoms which may be saturated or unsaturated. Examples of such oils are rapeseed oil, coriander oil, soyabean oil, cottonseed oil, sunflower oil, castor oil, olive oil, peanut oil, maize oil, almond oil, palm kernel oil, coconut oil and mustard seed oil. Rapeseed oil, which is a mixture of fatty acids partially esterified with glycerol, is preferred as it is available in large quantities and can be easily obtained by pressing from rapeseed.

Examples of derivatives of the fatty acids of vegetable or animal oils are alkyl esters, such as methyl esters. Such esters can be made by transesterification.

As lower alkyl esters of fatty acids, consideration may be given to the following, for example as commercial mixtures: the ethyl, propyl, butyl and especially methyl esters of fatty acids with 12 to 22 carbon atoms, for example of lauric acid, myristic acid, palmitic acid, palmitoleic acid, stearic acid, oleic acid, elaidic acid, 5 petroselic acid, ricinoleic acid, elaeostearic acid, linolenic acid, eicosanoic acid, gadoleic acid, docosanoic acid or erucic acid, which have an iodine number from 50 to 150, especially 90 to 125. Mixtures with particularly advantageous properties are those which contain mainly, ie, to at least 50 wt% methyl esters of fatty acids with 16 to 22 carbon atoms and 1, 2 or 3 double bonds. The preferred 10 lower alkyl esters of fatty acids are the methyl esters of oleic acid, linoleic acid, linolenic acid and erucic acid.

Commercial mixtures of the stated kind are obtained for example by cleavage and esterification of natural fats and oils by their transesterification with lower aliphatic 15 alcohols. For production of lower alkyl esters of fatty acids it is advantageous to start from fats and oils with high iodine number, such as, for example, sunflower oil, rapeseed oil, coriander oil, caster oil, soyabean oil, cottonseed oil, peanut oil or beef tallow. Lower alkyl esters of fatty acids based on a new variety of 20 rapeseed oil, the fatty acid component of which is derived to more than 80 wt% from unsaturated fatty acids with 18 carbon atoms, are preferred.

The hereinbefore described hydrocarbon oils may contain certain additives, depending on the intended use of the oil. For example, where the hydrocarbon oil 25 is a lubricating oil, it may contain viscosity index improvers such as ethylene-propylene copolymers, succinic acid based dispersants, metal containing dispersant additives and zinc dialkyl-dithiophosphate antiwear additives.

Where the hydrocarbon oil is a fuel oil, it may contain other additives such as 30 stabilisers, dispersants, antioxidants, corrosion inhibitors, cetane improvers and/or demulsifiers.

The mechanical system or device [of the third aspect of this invention]

Suitable as the mechanical system or device are those mechanical systems or 35 devices dependent for normal operation upon the flowability of hydrocarbon oil, particularly during periods where the oil temperature is below oil cloud point.

Typical of such mechanical systems are hydrocarbon oil storage- and distribution-systems, often being complex arrangements of vessels in liquid communication and dependent upon oil flowability for efficient oil transport around the system, usually by means of pumps. Such systems are typically found in refineries, oil 5 distribution terminals and networks, and on a smaller scale within appliances which utilise such oils, for example fuel oil installations and in-vehicle fuel oil systems. These mechanical systems typically contain devices such as filters and screens which strain insolubles from the oil and which are therefore themselves dependent, for normal operation, upon a continual through-flow of oil.

10 A decrease in oil flowability leads to a corresponding drop in oil passage through such systems and devices, reducing their efficiency of operation.

15 Using the oil compositions of this invention in such mechanical systems and devices, normal operation can continue at lower temperatures due to improved oil flowability, and particularly at lower temperatures below the oil composition cloud point.

20 However, such oil compositions may also be used at higher temperatures without detriment and provide assurance that, should oil composition temperature drop below cloud point, normal operation of the mechanical system or device will be maintained to lower temperatures.

25 The invention will now be illustrated by way of example only, as follows.

25 Examples of Comb Polymers

30 Comb polymers containing units of the formula (I) as hereinbefore described were prepared from the monomers in Table 1, using the standard polymerisation techniques also hereinbefore described.

Comparative polymers, being analogous to the above polymer examples but possessing larger hydrocarbyl substituents, were similarly prepared and are also shown in Table 1.

TABLE 1: COMB POLYMERS

Comb Polymer	1st Monomer	2nd Monomer	Molar Ratio of 1st:2nd monomer in polymer	Average carbon number of R11 substituent as measured over all units in the polymer	Average carbon number of R12 substituent as measured over all units in the polymer
Chemical Class	Carbon number of n-alkyl (ie. R11) substituents	Chemical class			
Polymer A	di-n-alkyl fumarate ester	8 and 10 in a 1:1 ratio	vinyl acetate	1:1	9
Polymer B	" 10 only	"	"	10	1
Polymer C	" 10 and 12 in a 1:1 ratio	"	"	11	1
Polymer D	" 10 and 12 in a 1:3 ratio	"	"	11.5	1
Comparative Polymer 1	" 12 only	"	"	12	1
Comparative Polymer 2	" 12 and 14 in a 1:1 ratio	"	"	13	1

Examples of Hydrocarbon Oils

The petroleum-based middle distillate fuel oils characterised in Table 2 were used to illustrate the invention.

5

TABLE 2: OIL CHARACTERISTICS

	Oils of the Invention				Comparative Oils	
	A	B	C	D	E	F
I.S.O. 3015 Cloud Point (°C)	-24	-27	-15	-14	-5	-4
E.N.116 (CFPP (°C))	-24	-27	-30	-29	-16	-16
ASTM D-86 Distillation						
IBP	182	190	157	140	162	171
20%	219	222	218	220	240	248
50%	248	240	259	267	283	268
90%	297	280	307	324	331	330
FBP	329	320	357	350	359	356
90%-20%	78	58	89	104	91	82
FBP-90%	32	40	50	26	28	26

Examples of the First and Second Aspects of the Invention

10

Oil compositions prepared by conventional blending techniques and illustrating the first aspect of the invention are defined in Table 3. The Cold Filter Plugging Point ('CFPP') of each composition was determined in accordance with the E.N. 116 standard test method, the CFPP values also being given in Table 3. The CFPP test is designed to correlate with the flow of a middle distillate fuel oil through the fuel systems of automotive diesels, at temperatures below oil composition cloud point. Fuel oils having greater flowability at such temperatures generally exhibit lower CFPPs.

15

20 In the CFPP test, a 40 ml sample of the oil is cooled in a bath maintained at about -34°C to give non-linear cooling at about 1°C/min.

Periodically, the cooled oil is tested for its ability to flow through a fine screen in a prescribed time period, using a test device comprising a pipette to whose lower end is attached an inverted funnel which is positioned below the surface of the oil.

5 Stretched across the mouth of the funnel is a 350 mesh screen having an area defined by a 12 millimetre diameter. Each periodic test is initiated by application of a vacuum to the upper end of the pipette, drawing oil through the screen up into the pipette until a mark indicating 20 ml of oil is reached. After each successful passage, the oil is returned immediately to the CFPP tube. The test is repeated

10 with each one degree drop in temperature until 20 ml of oil fails to pass through the screen within 60 seconds, the temperature at which failure occurs being reported as the CFPP temperature.

15 In Table 3, Co-additives I, II and III and IV are additional low temperature flow improvers suitable for use with the comb polymers of this invention.

20 Co-additive I is a polyoxyalkylene compound of the type hereinbefore described under linear compounds, being a behenic acid diester of a polyethylene glycol mixture predominant in glycols of molecular weights 200, 400 and 600.

25 Co-additive II is similar to co-additive I, being a mixed stearic acid/behenic acid diester of the same ethylene glycol mixture.

30 Co-additive III is a polar organic, nitrogen-containing compound of the class hereinbefore described, being the amide-amine salt formed by reacting one molar proportion of phthalic anhydride with two molar proportions of the secondary hydrogenated tallow amine, Armeen 2HT.

35 Co-additive IV is a conventional low temperature flow improver believed to be one or more ethylene vinyl-acetate or similar copolymers, but of unknown detailed composition and unknown treat rate ('x' in Table 3). Co-additive IV was already present in fuel oils C and D prior to testing, both these fuels having been purchased commercially.

TABLE 3

Example No.	Oil	Comb Polymer (ppm, a.i.)						Oil Composition				CFPP (°C)
		A	B	C	D	1	2	I	II	III	IV	
1.	A	-	-	-	-	-	-	-	-	-	-	-24
2.	A	70	-	-	-	-	-	280	-	-	-	-30.5
3.	A	-	70	-	-	-	-	280	-	-	-	-29.5
4.	A	-	-	70	-	-	-	280	-	-	-	-29.5
5.	A	-	-	-	70	-	-	280	-	-	-	-32
6.	A	-	-	-	-	70	-	280	-	-	-	-23
7.	B	-	-	-	-	-	-	-	100	-	-	-27
8.	B	-	25	-	25	-	-	-	100	-	-	-36
9.	B	-	-	-	-	25	-	-	100	-	-	-35
10.	B	-	-	-	-	-	25	-	100	-	-	-31
11.	B	-	-	-	-	-	-	25	-	100	-	-31
12.	B	-	-	-	-	-	-	-	-	-	-	-27
13.	C	-	-	-	-	-	-	-	-	-	x	-30
14.	C	-	120	-	-	120	-	-	-	-	120	-
15.	C	-	-	-	-	-	-	-	-	-	120	-
16.	C	-	-	-	-	-	-	-	-	-	120	-
17.	C	-	-	-	-	-	-	-	-	-	120	-
18.	E	-	-	-	-	-	-	-	-	-	-	-5
19.	E	-	150	-	-	-	-	-	-	-	150	-
20.	E	-	-	-	-	-	-	-	-	-	150	-
21.	F	-	-	-	-	-	-	-	-	-	150	-
22.	F	-	150	-	-	-	-	-	-	-	150	-
23.	F	-	-	-	-	-	-	-	-	-	-	-22

- means 'not present'

The results in Table 3 clearly illustrate the greater low temperature flowability of oil compositions in accordance with the first aspect of the invention. The oil compositions of examples 2 to 5 inclusive, 8 to 10 inclusive, 14 and 15 exhibit lower CFPPs than either the base fuels (examples 1, 7 and 13 respectively) or the oil compositions containing comparative comb polymers (examples 6, 11 and 12, 16 and 17 respectively).

The results in Table 3 similarly illustrate the second aspect of the invention. The CFPPs of oils having a cloud point no higher than -10°C (oils A, B and C) are effectively depressed by treatment with the comb polymers having hydrocarbyl substituents of average carbon number below 12, ie. Polymers A, B, C and D (examples 1 to 5 inclusive, 7 to 10 inclusive and 13 to 15 inclusive). In contrast, treatment of these oils with comparative polymers 1 or 2 has negligible effect upon the oil CFPP (examples 6, 11, 12, 16 and 17).

Similarly, comb polymers A to D inclusive prove less effective as CFPP depressants than comparative polymer 1 in fuels of cloud point higher than -10°C, ie. fuels E and F (examples 18 to 23 inclusive).

Oil compositions in accordance with this invention exhibit smaller wax crystals at temperatures below cloud point, consistent with their greater flowability at low temperatures. The oil composition defined in Table 4 were cooled from ambient temperature at 2°C per hour until -25°C was reached, where upon the wax crystals which had formed were photographed through an optical microscope.

25

TABLE 4

Example No.	Oil Composition				Crystal Picture
	Oil	Comb Polymer (ppm, ai)	Co-additive III (ppm, ai)		
26	D	180	—	—	180
27	D	—	180	—	180
28	D	—	—	180	180

--- means 'not present'

5 Examples 26 and 27 (oil compositions comprising comb polymers in accordance with this invention) clearly exhibit far smaller wax crystals at -25°C than example 28 (oil composition comprising a comb polymer not in accordance with this invention).

Example of the Third Aspect of the Invention

10 The CFPP results in Table 3 illustrate the greater flowability of oil compositions of the first aspect through a mechanical system comprising a fine screen (the CFPP testing apparatus).

15 The CFPP test was designed to correlate with the onset of automotive diesel engine system failure at low temperatures, this failure being due to fuel starvation resulting from the reduced flowability of the fuel oil through the vehicle fuel system. The lower CFPPs of the oil compositions of the first aspect thus indicate oil flowabilities sufficient to permit normal operation of such engine systems at lower temperatures, providing a technical advantage in regions of cold climate.

20 This advantage was confirmed by vehicle tests performed on a Cold Chamber Chassis Dynamometer, according to CEC Test Method M-11-T-91. In this test, a diesel-engined passenger car fuelled with test fuel oil is cooled in a cold climate chamber from 5°C above fuel oil cloud point to a temperature of -30°C over a 12 hour period, this latter temperature being held constant for 4 hours (a 'cold soak' period). The engine is then started from cold and the vehicle driven on a chassis dynamometer at a constant speed of 110 km/h, still at an air temperature of -30°C. The driveability performance of the vehicle is rated on a 'demerits' scale by an experienced operator where:

25 30 0 demerits corresponds to completely trouble-free driving, and 100 demerits corresponds to complete failure of the engine system.

Tests using a diesel-engined Ford Escort car were performed on the oil composition defined in Table 5, using additives hereinbefore described:

TABLE 5

Example No.	Oil Composition				Driveability Demerits at -30°C
	Oil	Comb Polymer B (ppm,ai)	Co-additive IV (ppm,ai)	Kerosene (%wt)	
24	D	180	x	-	0 demerits (PASS)
25	D	-	x	20%	48 demerits (FAIL)

'-' means not present

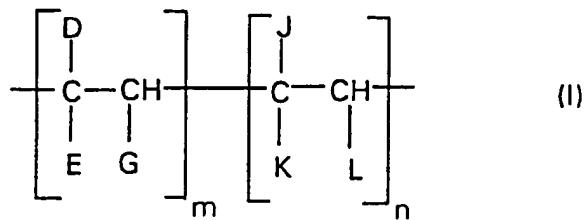
5

The addition of kerosene to a diesel fuel oil is common practice in cold regions, kerosene being a lighter petroleum fraction and serving to improve the cold temperature flowability of the diesel fuel oil. In this test the benefit from using the additive of this invention far outweighed that obtained by significant kerosene addition (20% wt, by weight of diesel fuel oil).

10

CLAIMS

1. An oil composition comprising a major proportion of hydrocarbon oil having a cloud point no higher than -10°C and a minor proportion of an additive comprising a comb polymer containing units of the general formula (I)



wherein D represents COOR¹¹, OCOR¹¹ or OR¹¹ groups,
 10 E represents H, CH₃, D or R¹² groups,
 G represents H or D group,
 J represents H, R¹² or an aryl or heterocyclic group,
 K represents H, COOR¹², OCOR¹², OR¹² or COOH groups,
 L represents H, R¹², COOR¹², OCOR¹² or aryl groups, and
 15
 wherein R¹¹ and R¹² each represent hydrocarbyl substituents of average carbon number below 12 as measured over all units in the polymer, and
 wherein m and n represent mole ratios, their sum being 1 and m being finite and being up to and including 1 and n being from zero to less than 1,
 20 provided that E, G, J, K and L do not each represent H when D represents COOR¹¹ or OCOR¹¹.

2. The oil composition of claim 1, wherein R¹² represents a hydrocarbyl substituent containing from 1 to 6 carbon atoms, and wherein R¹¹ represents a hydrocarbyl substituent different from R¹² and of average carbon number below 12.

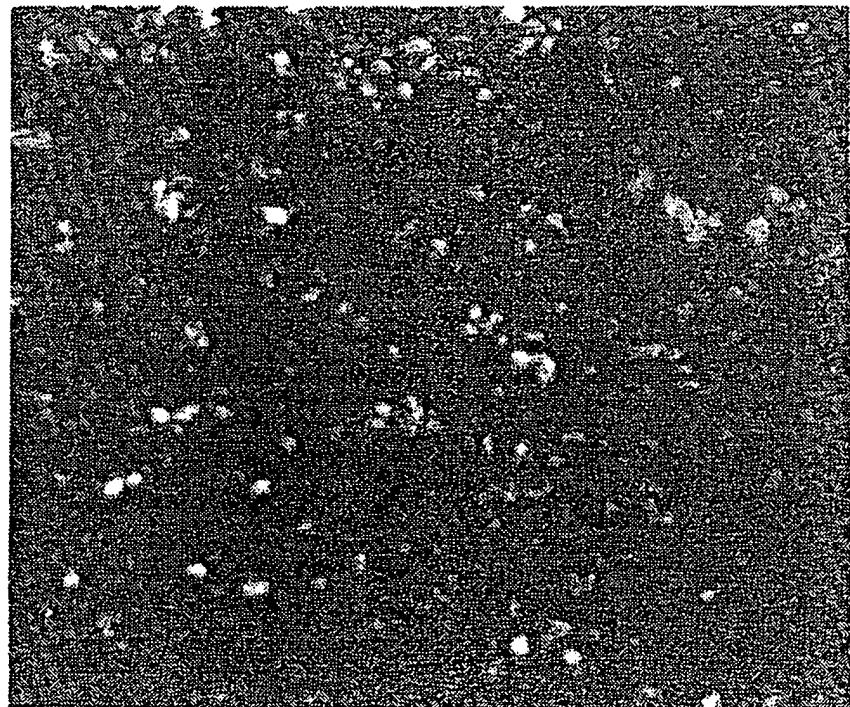
25
 3. The oil composition of claim 1 or claim 2 wherein the individual units of the polymer have R¹¹ substituents containing substantially the same number of carbon atoms.

30
 4. The oil composition of any preceding claim wherein the average carbon number of R¹¹ is at least 8.

5. The oil composition of any preceding claim wherein R¹¹ and R¹² are alkyl groups, and preferably n-alkyl groups.
6. The oil composition of any preceding claim wherein the hydrocarbon oil is a mineral fuel oil or a fuel oil derived from animal or vegetable material, or a mixture thereof.
7. The oil composition of any preceding claim wherein the hydrocarbon oil has a final boiling point no higher than 360°C and wherein the 20% and 90% distillation points of the hydrocarbon oil differ by less than 100°C.
8. The use of the additive defined in any of claims 1 to 5 for improving the low temperature flow properties of a hydrocarbon oil having a cloud point no higher than -10°C.
9. The use of the oil composition of any of claims 1 to 7 in a mechanical system or device dependent, for normal functioning, upon the flowability of hydrocarbon oil.
10. The use of claim 9, performed at temperatures below the oil composition cloud point.

FIG. 1

EXAMPLE 26



SUBSTITUTE SHEET (RULE 26)

FIG. 2

EXAMPLE 27



FIG. 3

EXAMPLE 28



INTERNATIONAL SEARCH REPORT

Int. Application No
PCT/EP 95/00666

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C10L1/18 C10M145/08 C10M145/16 C10M145/12 C10M145/02

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C10L C10M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB-A-573 364 (STANDARD OIL) 16 November 1945 see the whole document ---	1,3,5,6, 8-10
X	DE-A-21 02 469 (BASF) 3 August 1972 see the whole document ---	1,3,5,6, 8-10
X	WO-A-94 00537 (EXXON) 6 January 1994 see page 5 - page 6; claims 1-4,17 see page 17 ---	1,3-10
X	EP-A-0 214 786 (EXXON) 18 March 1987 see page 16, line 25 - line 38; claim 1 see page 19 ---	1,3-10
X	EP-A-0 225 688 (EXXON) 16 June 1987 see the whole document ---	1,3-10
		-/-

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *I* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

19 June 1995

Date of mailing of the international search report

21.7.95

Name and mailing address of the ISA

European Patent Office, P.O. 5818 Patenlaan 2
NL - 2280 HV Rijswijk
Tel. (+ 31-70) 340-2040, Tx. 31 651 cpo nl,
Fax (+ 31-70) 340-3016

Authorized officer

De La Morinerie, B

INTERNATIONAL SEARCH REPORT

International Application No PCT/EP 95/00666	
---	--

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
GB-A-573364				NONE
DE-A-2102469	03-08-72	BE-A-	778122	17-07-72
		FR-A, B	2122425	01-09-72
		GB-A-	1368159	25-09-74
WO-A-9400537	06-01-94	AU-B-	4561293	24-01-94
		CA-A-	2137229	06-01-94
		CN-A-	1081692	09-02-94
		EP-A-	0649456	26-04-95
EP-A-214786	18-03-87	CA-A-	1331511	23-08-94
		DE-A-	3686687	15-10-92
		JP-B-	7010983	08-02-95
		JP-A-	62096591	06-05-87
EP-A-225688	16-06-87	CA-A-	1277974	18-12-90
		DE-D-	3689374	20-01-94
		DE-T-	3689374	24-03-94
		JP-A-	62084186	17-04-87
FR-A-1534501		NONE		
GB-A-785570		NONE		
US-A-3961916	08-06-76	NONE		
WO-A-9101362	07-02-91	DE-A-	3924160	24-01-91
		AU-A-	6070890	22-02-91
		EP-A-	0483257	06-05-92
EP-A-217602	08-04-87	JP-A-	62070488	31-03-87
		US-A-	4802892	07-02-89
FR-A-982299	08-06-51	GB-A-	665130	
BE-A-658570	20-07-65	AU-A-	5322764	22-06-66
		FR-A-	1422550	16-03-66
		GB-A-	1080910	
		NL-A-	6500042	06-07-66

Siuda, Marcia

From: Jackson, Susan
Sent: Monday, August 26, 2002 11:11 AM
To: Siuda, Marcia
Subject: FW: 2000DE412 and 413



2000DE412



2000DE413



2000DE412

wly Revised Applicatent Application T. RACT OF THE DIS
Marcia

See me about this today for purposes of an IDS.

Susan

-----Original Message-----

From: Richard.Silverman@clariant.com
[mailto:Richard.Silverman@clariant.com]
Sent: Wednesday, July 10, 2002 12:25 PM
To: sjackson@kennedycoventon.com
Subject: 2000DE412 and 413

Here are copies of the subject cases which may be relevant to the prosecution of your 20010034968 A1. Both of the subject cases were filed in Germany on March 14, 2000 and filed in the US on March 14 2001. The serial numbers are 412: 09/808,454 and 413: 09/808,481.

Regards,

Dick

(See attached file: 2000DE412 Newly Revised Application Text.doc)(See attached file: 2000DE413 Patent Application Text.doc)(See attached file: 2000DE412 ABSTRACT OF THE DISCLOSURE.doc)